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Insights into Exo- and Endoglucanase Activities of Family 6 Glycoside Hydrolases from *Podospira anserina*

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The ascomycete *Podospira anserina* is a coprophilous fungus that grows at late stages on droppings of herbivores. Its genome encodes a large diversity of carbohydrate-active enzymes. Among them, four genes encode glycoside hydrolases from family 6 (GH6), the members of which comprise putative endoglucanases and exoglucanases, some of them exerting important functions for biomass degradation in fungi. Therefore, this family was selected for functional analysis. Three of the enzymes, *P. anserina* Cel6A (*PaCel6A*), *PaCel6B*, and *PaCel6C*, were functionally expressed in the yeast *Pichia pastoris*. All three GH6 enzymes hydrolyzed crystalline and amorphous cellulose but were inactive on hydroxyethyl cellulose, mannan, galactomannan, xyloglucan, arabinoxylan, arabinan, xylan, and pectin. *PaCel6A* had a catalytic efficiency on celotetraose comparable to that of *Trichoderma reesei* Cel6A (*TrCel6A*), but *PaCel6B* and *PaCel6C* were clearly less efficient. *PaCel6A* was the enzyme with the highest stability at 45°C, while *PaCel6C* was the least stable enzyme, losing more than 50% of its activity after incubation at temperatures above 30°C for 24 h. In contrast to *TrCel6A*, all three studied *P. anserina* GH6 cellulases were stable over a wide range of pHs and conserved high activity at pH values of up to 9. Each enzyme displayed a distinct substrate and product profile, highlighting different modes of action, with *PaCel6A* being the enzyme most similar to *TrCel6A*. *PaCel6B* was the only enzyme with higher specific activity on carboxymethylcellulose (CMC) than on Avicel and showed lower processivity than the others. Structural modeling predicts an open catalytic cleft, suggesting that *PaCel6B* is an endoglucanase.

Cellulose, a polysaccharide of β -1,4-linked D-glucose units, is the most abundant biopolymer on earth. It is the main constituent of plant cell walls, where it forms a tight complex together with hemicelluloses and is embedded in the lignin matrix. Cellulose is a recalcitrant material organized into microfibrils and composed of highly ordered glucan chains interlinked by hydrogen bonds. Depending on the source of cellulose, these fibrils have a more or less crystalline character, and for deconstruction of these complex structures, microorganisms have developed specialized enzymatic systems. All cellulolytic organisms produce multiple enzymes for cellulose degradation, but three main catalytic activities are necessary for complete hydrolysis: exoglucanases (or cellobiohydrolases [CBHs]) attack cellulose chains from the chain ends, and endoglucanases cleave the cellulose chain randomly, while β -glucosidases hydrolyze cellobiose, the reaction product of cellobiohydrolases. For efficient degradation, all three enzymatic activities have to be present, and synergistic interactions have been shown to be essential for a rapid degradation process (1–4). While exo-endo synergy can be easily explained by endoglucanases creating new chain ends for exoglucanases, exo-exo synergy still lacks a fully satisfactory explanation.

In contrast to synergistic interactions, the reaction mechanism of isolated enzymes is better understood, since structure-function studies have led to the identification of catalytic residues for cellobiohydrolases and endoglucanases (5–11). Sequence similarity and, therefore, structural properties are at the origin of the classification of these enzymes in the CAZY (Carbohydrate-Active enzymes) database, facilitating the assignment of function (12). Fungal endoglucanases are classified into glycoside hydrolase families GH5, GH6, GH7, GH9, GH12, GH45, and GH74, while fungal exoglucanases can be found in families GH6, GH7, and GH48.

At the three-dimensional (3D) level, cellobiohydrolases are

characterized by a typical tunnel-shaped active site, with the roof of the tunnel formed by long flexible loops (5, 13–15). In contrast, their homologous endoglucanases bear shorter loops converting their active site into a classical cleft. This difference in active-site topology has consequences in the action pattern of the enzymes. The cellulose chain end is believed to enter the tunnel active site of cellobiohydrolases and to be cut processively into cellobiose units as it threads through the tunnel (5, 16). Instead, the open cleft active site of endoglucanases is thought to allow random binding and cleavage along the cellulose chain.

An increasing number of genomic and proteomic studies on fungi reveal a large variety of lignocellulose-degrading enzymes present in these organisms. A very interesting ascomycete is *Podospira anserina*, which grows on herbivore dung. Its mycelium develops at a late stage, after the most easily utilizable biomass components, such as hemicellulose and pectin, have already been degraded by other species. *P. anserina* is therefore hypothesized to be able to specifically use the recalcitrant parts of lignocellulose. The analysis of its genome has indeed revealed an impressive number of genes encoding carbohydrate-activating enzymes (CAZymes) and enzymes putatively involved in lignin degradation, including oxidoreductases, cellobiose dehydrogenase, cop-

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per radical oxidases, or laccases, which is a rather atypical feature for ascomycetes (17). The total number of putative glycoside hydrolases is similar to those in other sequenced ascomycetes, but it has the largest panel of cellulose-degrading enzymes and carbohydrate binding modules (CBMs). For instance, *P. anserina* has 30 genes encoding putative cellulases belonging to the families cited above, compared to only 8 for *Trichoderma reesei*, a number which is particularly small compared to most other carbohydrate-degrading ascomycetes (18).

The specific activity of CBHs on soluble and insoluble substrates has been shown to be lower than that of endoglucanases (19), and the CBH activity is probably rate limiting for cellulose hydrolysis (20). For efficient hydrolysis, a large amount of CBH enzyme is therefore needed in cellulolytic complexes, such as the one of *T. reesei*. In the latter organism, CBHs are the most abundant enzymes; i.e., Cel7A (CBH1) makes up 40 to 60% and Cel6A (CBH2) makes up about 20 to 30% of the total amount of secreted proteins (21). An enhancement of Cel6A was shown to be beneficial for hydrolytic activity of the complex (22), and Cel6A deletion mutants showed a 33% decrease in saccharification efficiency (23). Intriguingly, *T. reesei* contains only one enzyme of the GH6 family, in contrast to *Humicola insolens* and *Myceliophthora thermophila*, two other fungi that produce efficient cellulolytic cocktails containing three family GH6 enzymes. *H. insolens* Cel6A and Cel6B have been shown to display endoglucanase and cellobiohydrolase activities, respectively (24–26). *P. anserina* has four genes encoding putative GH6 enzymes, but to date, nothing is known about the enzymatic properties of this important family of cellulose-degrading enzymes in this organism. The reason for the existence of multiple enzymes is not known, and the question arises as to whether there is redundancy or if each enzyme has a distinct role.

To answer this question, we undertook the cloning of the four *P. anserina* genes encoding putative GH6 cellulases to express them heterologously. We successfully produced three of the enzymes, termed *P. anserina* Cel6A (*PaCel6A*), *PaCel6B*, and *PaCel6C*. Substrate and product profiles of the purified enzymes revealed significant differences concerning the specificities and modes of action on cellulose model substrates. The biochemical characteristics of *P. anserina* GH6 cellulases were compared to those of *T. reesei* Cel6A (*TrCel6A*), revealing differences in their modes of action.

MATERIALS AND METHODS

Culture media. *P. anserina* was grown at 27°C on M2 medium, composed of 0.25 g liter⁻¹ KH₂PO₄, 0.3 g liter⁻¹ K₂HPO₄, 0.25 g liter⁻¹ MgSO₄ · 7H₂O, 0.5 g liter⁻¹ urea, 0.05 g liter⁻¹ thiamine, 0.25 g liter⁻¹ biotin, 2.5 mg liter⁻¹ citric acid, 2.5 mg liter⁻¹ ZnSO₄, 0.5 mg liter⁻¹ CuSO₄, 125 μg liter⁻¹ MnSO₄, 25 μg liter⁻¹ boric acid, 25 μg liter⁻¹ Na₂MoO₄, 25 μg liter⁻¹ iron alum, 5 g liter⁻¹ dextrin, and 10 g liter⁻¹ yeast extract, adjusted to pH 7.0 with KH₂PO₄. For *Pichia pastoris*, three media were used: minimum methanol (MM), composed of 3.4 g liter⁻¹ yeast nitrogen base (YNB) (Difco), 10 g liter⁻¹ ammonium sulfate, 20 g liter⁻¹ agar, 2 ml of 200 g liter⁻¹ biotin, and 5 ml of pure methanol; BMGY, containing 3.4 g liter⁻¹ YNB, 10 g liter⁻¹ ammonium sulfate, 10 g liter⁻¹ glycerol, 10 g liter⁻¹ yeast extract, 10 g liter⁻¹ peptone, 100 ml of 1 M phosphate buffer (pH 6.0), and 2 ml of 200 g liter⁻¹ biotin; and BMMY, which is identical to BMGY except that it contains 30 ml liter⁻¹ of pure methanol instead of glycerol.

Cloning procedures. *P. anserina* strain S mat+, which was used in this study, was kindly provided by P. Silar (CNRS, Paris, France). *P. anserina*

cells were grown in baffled flasks at 120 rpm and at 27°C on M2 medium (0.25 g liter⁻¹ KH₂PO₄, 0.3 g liter⁻¹ K₂HPO₄, 0.25 g liter⁻¹ MgSO₄ · 7H₂O, 0.5 g liter⁻¹ urea, 0.05 mg liter⁻¹ thiamine, 0.25 μg liter⁻¹ biotin, 2.5 mg liter⁻¹ citric acid, 2.5 mg liter⁻¹ ZnSO₄, 0.5 mg liter⁻¹ CuSO₄, 125 μg liter⁻¹ MnSO₄, 25 μg liter⁻¹ boric acid, 25 μg liter⁻¹ Na₂MoO₄, 25 μg liter⁻¹ iron alum, 5 g liter⁻¹ dextrin [pH 7]) supplemented with 1% Avicel cellulose, with or without induction by 0.1% sophorose 1 h prior to harvesting of the mycelia. Total RNA was extracted from 3- or 5-day-old cultures with the RNeasy plant kit (Qiagen, Courtaboeuf, France), and cDNAs were synthesized by using SuperScript reverse transcriptase (RT) (Life Technologies, NY, USA), according to the manufacturer's instructions. *PaCel6B* and *PaCel6D* were amplified by PCR using *Pfu* DNA polymerase (Promega, WI, USA) and the following primers: *PaCel6B*-F (5'-TAG AAT TCG CCC CTT CCC CGA CCA CC-3'), *PaCel6B*-R (5'-GAT CTA GAC CGA GAA GGG AAG GGT TAG A-3'), *PaCel6D*-F (5'-TAG AAT TCT CTC CCC TTG AGG CAC GC-3'), and *PaCel6D*-R (5'-GAT CTA GAC CCA AGC ACT GCG AAT ACC A-3'). *PaCel6A* and *PaCel6C* could not be obtained by amplification, and their coding sequences were synthesized after codon optimization for expression in *P. pastoris* (Eurogentec, Belgium). The *TrCel6A* gene was amplified from cDNA obtained from *T. reesei* strain CL847. Coding sequences for *PaCel6A*, *PaCel6B*, *PaCel6C*, and *TrCel6A* were cloned into the pPICZαA vector (Life Technologies), in frame with the yeast α-secretion factor and with C-terminal hemagglutinin (HA) and His tags. For an unknown reason and despite multiple trials, the *PaCel6D* coding sequence, which was obtained after RT-PCR, could not be cloned into the expression vector and therefore could not be produced. The enzyme is therefore termed *PaGH6D*.

Competent X33 yeast cells were prepared and transformed by electroporation according to the EasySelect *Pichia* expression kit protocol (Life Technologies). Expression vectors were linearized by *Pme*I (New England BioLabs, MA, USA) prior to transformation. After electroporation, cells were spread onto yeast extract-peptone-dextrose medium with sorbitol (YPDS)-zeocin plates. Transformants were selected by their lower growth rates on MM plates than on minimum dextrose (MD) plates.

Enzyme production and purification in *Pichia pastoris*. Culture supernatants of 10 clones were first analyzed by small-scale protein productions in a total volume of 10 ml BMGY, as described previously (27). When the optical density at 600 nm (OD₆₀₀) reached a value of between 2 and 6, cells were transferred into 2 ml BMMY and grown at 30°C, with daily addition of 3% (vol/vol) methanol. After SDS-PAGE analysis, clones with the highest secretion levels were selected, cultured in 200 ml BMGY starting culture, and transferred into 40 ml BMMY. Recombinant proteins were recovered after 5 days of methanol induction. After 10 min of centrifugation at 4,000 × g, supernatants were passed through a 0.2-μm filter. Samples were then concentrated 10 times in binding buffer (50 mM Tris-HCl [pH 7.8], 150 mM NaCl, 10 mM imidazole) by ultrafiltration in a Vivaspin 20 column (polyethersulfone membrane, 10-kDa cutoff; Sartorius, France). Subsequently, recombinant proteins were purified on a 5-ml HisTrap column (GE Healthcare) connected to an Äkta fast protein liquid chromatography (FPLC) apparatus (GE Healthcare), according to the manufacturer's instructions. Enzymes were eluted with binding buffer supplemented with 150 mM imidazole. A final ultrafiltration step was used to concentrate proteins in 50 mM acetate buffer (pH 5). Protein homogeneity was checked on a 12% SDS-polyacrylamide gel, followed by Coomassie staining.

Protein analysis. Deglycosylation of 2 μg recombinant protein was performed with 2 μl of either endo-α-N-acetylgalactosaminidase and 2 μl neuraminidase or 2 μl endoglycosidase H (Endo H) (all from New England BioLabs) for 2 h at 37°C, according to the manufacturer's instructions.

N-terminal sequences were determined by Edman degradation with a Procise cLC sequencing system (model 494cLC; Applied Biosystems) from purified protein samples electroblotted onto a polyvinylidene difluoride (PVDF) membrane (Life Technologies).

TABLE 1 Molecular characteristics of the four *P. anserina* GH6 enzymes and identities with known enzymes^a

Enzyme	Gene ID ^b	Molecular mass (kDa) (pI) of encoded protein	CBM positions	Linker positions	Catalytic domain positions	No. of predicted glycosylation sites		% identity to <i>Tr</i> Cel6A (CBH2) ^c	Database entry with highest similarity, GenBank accession no. (% similarity)
						N-Glycosylation	O-Glycosylation		
<i>Pa</i> Cel6A	6187194	49.3 (5.8)	9–45	46–102	103–467	1	29	67.8	<i>Chaetomium globosum</i> hypothetical protein, EAQ82944 (77)
<i>Pa</i> Cel6B	6187311	41.3 (6.2)			1–364	3	2	35.4	<i>Myceliophthora thermophila</i> GH6, AEO57190 (73)
<i>Pa</i> Cel6C	6188027	40.9 (5.1)			1–380	1	3	45.8	<i>Sordaria macrospora</i> hypothetical protein, XP_003344888 (79)
<i>Pa</i> GH6D	6187939	44.0 (7.5)	377–412	349–376	1–348	1	9	35.0	<i>Humicola insolens</i> endoglucanase 6B, Q7SIG5 (82)

^a Amino acid ranges of functional domains of mature proteins were predicted by InterProScan.

^b Gene identifier as in the *P. anserina* reference genome sequence.

^c Identity calculated with catalytic domains only.

Biochemical characterization by activity assays. Determinations of activity profiles on solid substrates were performed with microtiter plates by measuring the release of reducing sugars with 3,5-dinitrosalicylic acid (DNS), as reported previously (28). The total volume was 100 μ l, containing 6.4 mg liter⁻¹ of purified enzymes in 50 mM acetate buffer (pH 5) for *Tr*Cel6A or citrate-phosphate buffer (pH 7) for the three *Pa*Cel6 enzymes and 10 g liter⁻¹ substrate. Substrates used were Avicel PH-101 cellulose, carboxymethylcellulose (CMC), hydroxyethyl cellulose (HEC), birchwood xylan, pectin (all from Sigma-Aldrich), barley β -glucan, 1,4- β -D-mannan, carob galactomannan, sugar beet arabinan, wheat arabinoxylan, arabinogalactan, xyloglucan, and konjak glucomannan (all from Megazyme, Wicklow, Ireland). Reaction mixtures were incubated at 35°C or 45°C, depending on enzyme stability, for 30 min. Reactions were stopped by boiling, and the mixtures were centrifuged at 4,000 \times g for 3 min.

Activities on soluble cellooligosaccharides were measured at optimal pH and temperature with 20 mg/liter cellooligosaccharides and 10 mg liter⁻¹ enzyme. Reactions were stopped after 30 min by placing the mixtures into boiling water for 5 min. Reaction products were analyzed with a Dionex ISC300 high-performance liquid chromatography (HPLC) system as described previously (29). For determination of kinetic constants, enzymes were incubated at an appropriate concentration (0.5 to 2 nM) in 50 mM phosphate buffer (pH 7) and at optimal temperature (45°C for *Pa*Cel6A and *Pa*Cel6B and 35°C for *Pa*Cel6C) with cellotetraose (0.5 to 30 μ M) for up to 20 min. Reactions were inactivated by placing the mixtures into a boiling water bath for 5 min, and reaction products were analyzed by HPEAC-PAD (high-performance anion-exchange chromatography coupled with pulsed amperometric detection). Kinetic constants were determined by using the least-squares method.

Product profiles were established in duplicates on Avicel and CMC at 1% dry matter in 10 mM citrate-phosphate buffers. The substrates were incubated with 10 mg g⁻¹ enzyme for 15 min and 24 h under optimal conditions (*Tr*Cel6A, 45°C at pH 5; *Pa*Cel6A, 45°C at pH 7; *Pa*Cel6B, 35°C at pH 7; *Pa*Cel6C, 25°C at pH 6). Reactions were stopped by boiling for 5 min, and reaction products (glucose, cellobiose, and cellotriose) were analyzed by HPEAC-PAD. Processivity was determined by using oligosaccharide ratios, as described previously (30, 31).

Effect of pH and temperature on enzymatic activity. The apparent optimal pH was estimated by using 1% Avicel in a total volume of 100 μ l of 50 mM citrate-phosphate buffer (pH 3 to 7), 50 mM Tris-HCl (pH 7 to 9), and 50 mM Tris-maleate (pH 9 to 10). The optimal reaction temperature was studied by incubating the enzymes with 1% Avicel for 15 min at temperatures of between 30°C and 70°C, at pH 5 for *Tr*Cel6A and at pH 7 for *Pa*Cel6 enzymes. For enzymatic stabilities, enzymes were incubated in different buffers or at different temperatures for 24 h, and residual activity was measured as described above. Reducing sugars were determined according to a protocol adapted from a method described previously (32). Briefly, 40 μ l of supernatant or glucose standard was added to 120 μ l of reagent 1, made extemporarily from 2 volumes of reagent 1A [0.5 g liter⁻¹ K₃Fe(CN)₆, 34.8 g liter⁻¹ K₂HPO₄ (pH 10.6)] and 1 volume of reagent 1B

(5.3 g liter⁻¹ Na₂CO₃ and 0.65 g liter⁻¹ KCN). The plates were sealed and heated to 96°C. After 10 min at room temperature, 80 μ l of the reaction mix was added to 40 μ l of reagent C (2.5 g liter⁻¹ FeCl₃, 20 g liter⁻¹ polyvinyl pyrrolidone K25, 2 N H₂SO₄). The optical density was measured at 520 nm after 15 min of incubation at room temperature in the dark. Each point was the mean of three independent measurements, with a standard deviation of 2 to 10%.

Cellulose binding assay. The capacity of adsorption of enzymes to Avicel was determined after incubating 60 μ g of protein with 300 μ l 1% Avicel PH-101 (Sigma-Aldrich) in 50 mM acetate buffer (pH 5) for *Tr*Cel6A or citrate-phosphate buffer (pH 7) for the three *Pa*Cel6 enzymes for 4 h at 4°C with agitation. After centrifugation, the amount of cellulases left in the supernatant was quantified by using the Bio-Rad protein assay kit with bovine serum albumin (BSA) as a standard (Bio-Rad, France). Controls without protein or with BSA were analyzed in parallel.

Structure modeling and bioinformatic analysis. Sequence alignments were done by using ClustalW on the EMBL-EBI server (<http://www.ebi.ac.uk/Tools/msa/clustalw2/>). *Pa*Cel6 structures were predicted by using the PHYRE2 server (<http://www.sbg.bio.ic.ac.uk/phyre2/>) (33). Models were visualized with PyMOL (PyMOL Molecular Graphics System, version 1.1; DeLano Scientific, USA). All *Pa*Cel6 models were registered in the Protein Model Database (<http://www.caspar.it/PMDB/>). PMDB identifiers are PM0077882, PM0077884, PM0077885, and PM0077886 for *Pa*Cel6A, *Pa*Cel6B, *Pa*Cel6C, and *Pa*Cel6D, respectively.

The NetOGlyc (<http://www.cbs.dtu.dk/services/NetOGlyc/>) and NetNGlyc (<http://www.cbs.dtu.dk/services/NetNGlyc/>) servers were used for glycosylation site predictions.

RESULTS

Sequence analysis and expression of *P. anserina* GH6 enzymes. The annotated *P. anserina* genome has four genes encoding family GH6 glycoside hydrolases (GenBank accession numbers XM_001903135 for *Pa*Cel6A, XM_001903174 for *Pa*Cel6B, XM_001903858 for *Pa*Cel6C, and XM_001903610 for *Pa*GH6D). *Pa*Cel6A and *Pa*GH6D both harbor a cellulose binding module of family CBM1 at their N terminus, explaining their slightly higher molecular masses (Table 1). *Pa*Cel6B, *Pa*Cel6C, and *Pa*GH6D catalytic domains have very limited identity to the *T. reesei* cellobiohydrolase 2 enzyme (*Tr*Cel6A).

At the time of writing, the most similar proteins that have been biochemically characterized are exoglucanase A from *Humicola insolens* (UniProtKB accession number Q9C1S9), displaying 72% and 50% identities to *Pa*Cel6A and *Pa*Cel6C, respectively, and *H. insolens* endoglucanase 6B (accession number Q7SIG5), with 54 and 82% identities to *Pa*Cel6B and *Pa*GH6D, respectively. Identities between the four *Pa*Cel6 enzymes ranged between 25 and 45%, with *Pa*Cel6B and *Pa*GH6D displaying the highest similarity

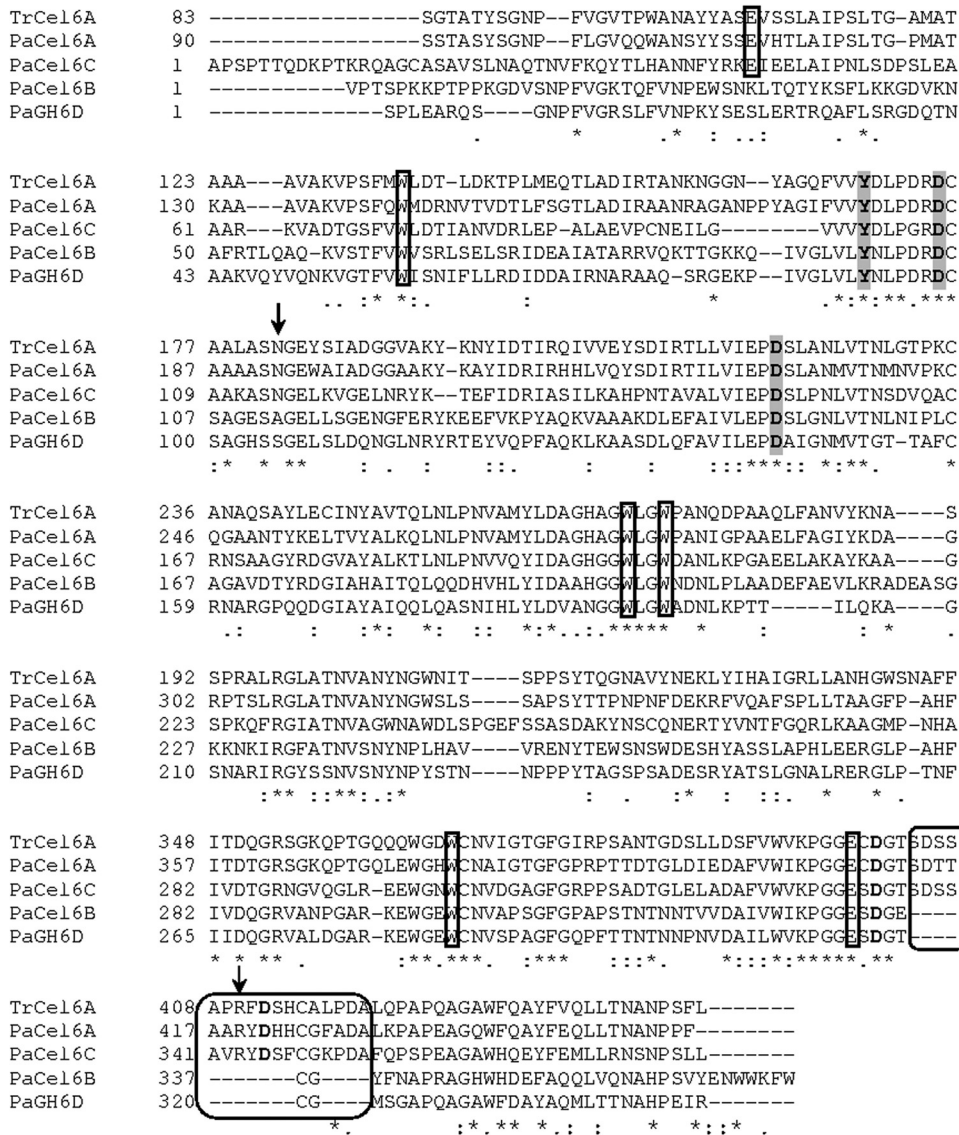


FIG 1 Amino acid sequence alignment of catalytic domains of *T. reesei* Cel6A and the four *P. anserina* family 6 glycoside hydrolases. Conserved residues are marked by an asterisk. Amino acids involved in catalysis are shaded. Amino acids implicated in glucose binding are boxed. Residues N182 and R410, which establish a direct contact between the two surface loops in *TrCel6A*, are indicated by an arrow. The sequence stretch corresponding to the second surface loop present in *TrCel6A*, *PaCel6A*, and *PaCel6C*, but absent in *PaCel6B* and *PaGH6D*, is boxed.

to each other and *PaCel6A* and *PaGH6D* displaying the lowest similarity.

Sequence analysis of the four *PaCel6* enzymes was realized by a multiple alignment including *TrCel6A* (Fig. 1). All amino acids which are known to be involved in catalysis, namely, Asp221 (following *TrCel6A* numbering) and Asp175 (7), are conserved in all four *PaCel6* sequences. Tyr169, proposed to participate in catalysis by causing distortion of the glucose ring undergoing catalysis (34), was also found to be conserved in all four proteins. Two other aspartic acid residues, Asp401 and Asp412, not directly involved in catalysis but contributing to full enzymatic activity (11), are conserved in *PaCel6A* and *PaCel6C*, but Asp412 is absent in *PaCel6B* and *PaGH6D*. The most important amino acids involved in binding of the glucose moiety at the -2, +1, +2, and +4 subsites (throughout this paper, we follow the subsite nomencla-

ture proposed by Davies et al. [35]) are also present in all four *PaCel6* enzymes (Trp135, Trp269, Trp272, and Trp367) (9), but Glu107 (-3 subsite) is not conserved in *PaCel6B* and *PaGH6D* (11).

In order to induce the expression of *PaCel6* genes, *P. anserina* was grown on Avicel as the sole carbon source. Under this condition, only cDNAs encoding *PaCel6B* and *PaGH6D* were obtained, whereas synthetic genes had to be generated for cloning of *PaCel6A* and *PaCel6C* in *P. pastoris*. For an unknown reason, cloning of the *PaGH6D* coding sequence failed despite several trials, and the corresponding protein therefore could not be produced in *P. pastoris*. The three other proteins, *PaCel6A*, *PaCel6B*, and *PaCel6C*, finally yielded 30 to 130 mg liter⁻¹ of protein after induction of *P. pastoris* transformants by methanol. In parallel, *TrCel6A* was also expressed in *P. pastoris* for comparative studies.

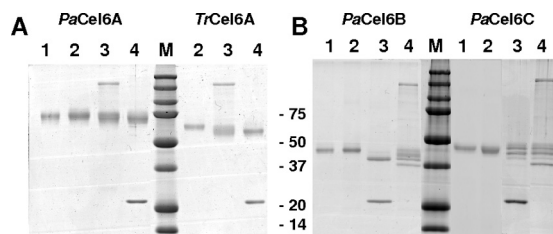


FIG 2 SDS-PAGE of Ni-purified recombinant glucanases. (A) *PaCel6A* and *TrCel6A*. Lanes 1 and 2, recombinant proteins; lanes 3, treatment with endo- α -N-acetylgalactosaminidase (O-glycosidase); lanes 4, treatment with Endo H. (B) *PaCel6B* and *PaCel6C*. Lanes 1 and 2, recombinant proteins; lanes 3, treatment with Endo H; lanes 4, treatment with endo- α -N-acetylgalactosaminidase (O-glycosidase). M, molecular mass marker (in kilodaltons) (Dual Precision Plus; Bio-Rad).

After purification on a Ni column, proteins were analyzed by SDS-PAGE (Fig. 2). All proteins displayed higher apparent molecular masses than the predicted ones, suggesting glycosylation by the host. Whereas *PaCel6B* and *PaCel6C* seemed only slightly glycosylated, *PaCel6A* and *TrCel6A* had an apparent molecular mass of about 70 kDa (theoretical mass of *TrCel6A* of 47.9 kDa). For all proteins, both N- and O-glycosylation sites are predicted (Table 1). Deglycosylation by Endo H led to reductions of the molecular masses of all proteins. The most important reduction was observed for *TrCel6A* and *PaCel6B*, which also have the largest number of predicted N-glycosylation sites (Fig. 2). A large number of O-glycosylation sites are predicted for the linker domain of *TrCel6A* and *PaCel6A* (Table 1). Treatment with endo- α -N-acetylgalactosaminidase reduced the molecular masses of all recombinant proteins. Several bands appeared after deglycosylation of *PaCel6B* and *PaCel6C*, which is supposed to be due to incomplete deglycosylation. The molecular mass of *PaCel6A* could not be reduced significantly by either N- or O-deglycosylation. A lower level of incorporation of SDS or inefficient deglycosylation could explain the migration at a higher apparent molecular mass.

The N-terminal sequence was determined for all proteins, which confirmed that all of them were correctly processed, except for *PaCel6A*, which, for an unknown reason, lacked the first 5 amino acids. The N-terminal sequence of the mature protein started with ERQN. However, this should not impact the function of the CBM, which is situated at the N terminus, but is predicted to start only 8 amino acids further.

Enzymatic properties of *P. anserina* GH6 enzymes. *PaCel6* enzymatic activities were assayed on celooligosaccharides and structurally different polysaccharides. Table 2 summarizes the product profiles on G3 (celotriose) to G6 (cellohexaose) cellooligosaccharides. The observed cleavage pattern for *TrCel6A* confirmed previous results for the native enzyme (36). *PaCel6* enzymes all had similar cleavage patterns, with the exception that glucose was never detected after cleavage by *PaCel6A*, in contrast to the three other enzymes. All enzymes were active on Avicel and CMC. In addition, *PaCel6B* displayed activity on barley β -(1,3;1,4)-glucan and glucomannan. No activity was detected on mannan, galactomannan, arabinoxylan, arabinan, xylan, xyloglucan, HEC, and pectin. As all enzymes were active on Avicel, this substrate was chosen to determine the apparent temperature and pH optima as well as stability. *PaCel6* enzymes had rather narrow temperature optima, which were generally lower than that of *TrCel6A*. *PaCel6A* presented the highest activity at 55°C but had a 30%

lower activity at 65°C, in contrast to *TrCel6A*, which displayed the same activity at both temperatures (Fig. 3A). Temperature stability profiles showed that *PaCel6A* was the most stable of the *P. anserina* enzymes (up to 45°C, as for *TrCel6A*), whereas *PaCel6B* and *PaCel6C* lost nearly all activity after 24 h at 45°C and 35°C, respectively (Fig. 3B).

The Cel6 enzymes from *P. anserina* showed interesting activity profiles when the pH varied (Fig. 3C). These enzymes maintained nearly 100% of their activity from pH 5 up to pH 9, in contrast to *TrCel6A*, which lost about 50% of its activity at pH ≥ 6 . Concerning stability, *PaCel6A* was also fairly stable for 24 h at these pH values. The activities of *PaCel6B* and *PaCel6C* declined upon incubation for 24 h at pH ≥ 7 . The following experiments including the determination of kinetic parameters were conducted under optimal conditions for each enzyme.

Michaelis-Menten constants were determined on the soluble substrate cellotetraose. This substrate was cleaved by all four enzymes exclusively into two cellobiose units. HPAEC-PAD analyses revealed only traces of glucose and celotriose. The kinetic behavior was very different for each *P. anserina* enzyme (Table 3). Kinetic constants for *PaCel6A* and *TrCel6A* were rather similar. The k_{cat} value of *PaCel6A* was 2-fold higher than that of *TrCel6A*, but it was determined at the optimal temperature for this enzyme (45°C), in contrast to *TrCel6A* (27°C). Interestingly, *PaCel6B* had a very high turnover number ($k_{cat} = 27.7 \text{ s}^{-1}$) but the lowest affinity for cellotetraose ($K_m = 43 \text{ }\mu\text{M}$). The k_{cat} of *PaCel6C* was about 100-fold lower than that of *PaCel6B*, but the affinity for cellotetraose was much higher. The catalytic efficiencies (k_{cat}/K_m) of *PaCel6B* and *PaCel6C* were 2-fold lower than those of *PaCel6A* and *TrCel6A*.

Product profile analyses. Because glycoside hydrolase family GH6 groups together both exo- and endoglucanases, we have determined the product profiles of the three *P. anserina* GH6 enzymes, using CMC and Avicel as substrates (Table 4). The main hydrolysis product from these cellulose substrates was cellobiose (G2) and, to a lesser extent, glucose (G1) and celotriose (G3). The fact that *TrCel6A* slowly cleaves G3 leads to a larger amount of G3

TABLE 2 Product profile of recombinant *PaCel6* enzymes and *TrCel6A* on soluble oligosaccharides

Enzyme	Substrate	Product(s)
<i>TrCel6A</i>	G3	G1, G2, G3
	G4	G2
	G5	G1, G2, G3
	G6	G1, G2, G3
<i>PaCel6A</i>	G3	G3
	G4	G2
	G5	G2, G3
	G6	G2, G3
<i>PaCel6B</i>	G3	G1, G2, G3
	G4	G2
	G5	G2, G3
	G6	G1, G2, G3
<i>PaCel6C</i>	G3	G1, G2, G3
	G4	G2
	G5	G2, G3
	G6	G2, G3

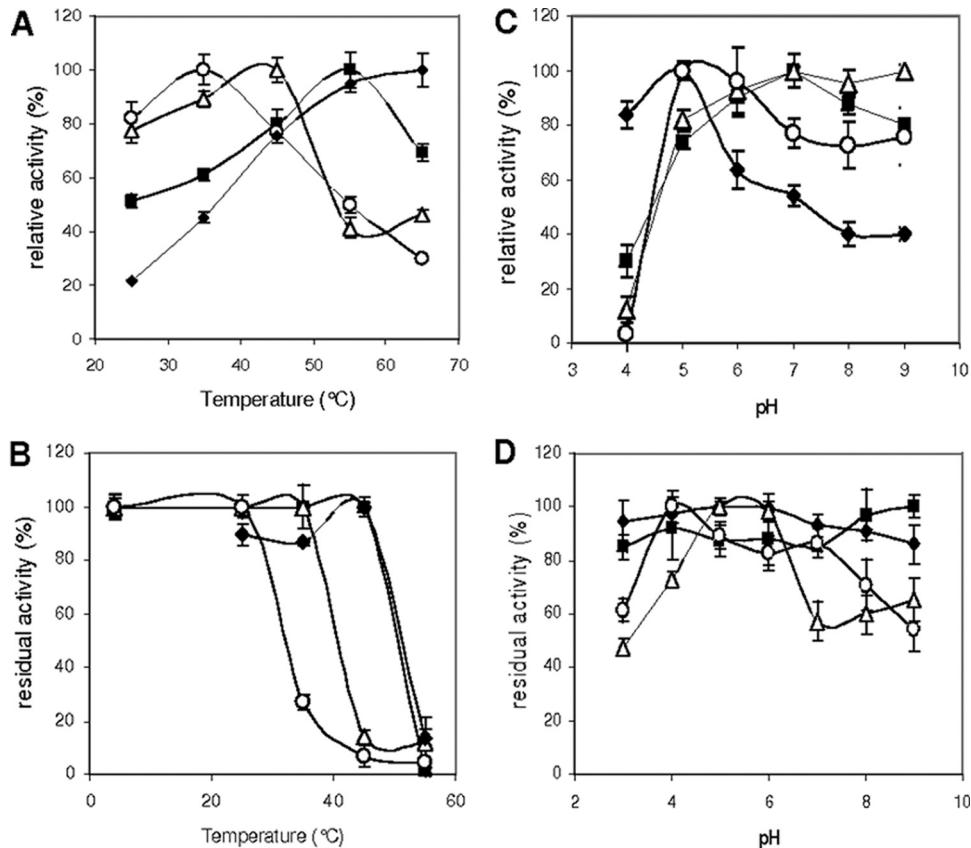


FIG 3 Activity of *P. anserina* Cel6 and *T. reesei* Cel6A on Avicel as a function of temperature and pH. (A) Optimal temperature. Avicel (1%) was hydrolyzed for 15 min at the indicated temperatures. (B) Temperature stability. Activity was measured after incubation at the indicated temperatures for 24 h. Hydrolysis reactions were conducted at 35°C. (C) Optimal pH. Hydrolysis reactions were conducted at 35°C for 15 min. (D) pH stability. Activity was measured at 35°C after incubation at the indicated pHs for 24 h. Symbols: filled diamonds, *TrCel6A*; filled squares, *PaCel6A*; open triangles, *PaCel6B*; open circles, *PaCel6C*. Values are means of triplicate measurements.

and lower G2/G3 ratios at 15 min than at 24 h. The G2/G1 ratio decreased slightly or remained the same at 24 h, which is in accordance with the formation of G1 following the cleavage of G3.

For *PaCel6A*, the G3 concentration increased from 6 to 49 μM between 15 min and 24 h, but the G2/G3 ratio increased at the same time. A similar trend was observed for G1; i.e., the glucose amounts and the G2/G1 ratio increased with time. These data suggest that G3 and G1 might be formed mainly in the first minutes of the reaction, by an initial attack liberating G1 or G3 instead of the G2 produced in a processive action (“false initial attack”

TABLE 3 Specific activities of recombinant enzymes on cellulose substrates and kinetic constants for *PaCel6A*, *PaCel6B*, and *PaCel6C*, compared to the constants for *TrCel6A*, measured on cellotetraose under optimal conditions for each enzyme

Enzyme	Sp act (nmol min ⁻¹ mg ⁻¹)		K_m (μM)	k_{cat} (s ⁻¹)	k_{cat}/K_m (μM^{-1} s ⁻¹)
	Avicel	CMC			
<i>TrCel6A</i> ^a	16.4	6.7	2.6	3.1	1.2
<i>PaCel6A</i>	7.6	3.0	4.7	6.6	1.4
<i>PaCel6B</i>	2.3	5.5	43.0	27.7	0.65
<i>PaCel6C</i>	1.9	1.1	0.59	0.3	0.54

^a Kinetic constants reported previously (36), measured at 27°C and pH 5.

TABLE 4 Hydrolysis products from 1% Avicel or CMC with *P. anserina* GH6 enzymes or *TrCel6A*, measured after 15 min (Avicel) and 24 h (Avicel and CMC) of hydrolysis under optimal conditions

Treatment and enzyme	Concn (μM) ^a				Processivity ^b
	Glucose	Cellulose	Cellotriose	Total	
Avicel 15 min					
<i>TrCel6A</i>	9.2	112	5.7	127	
<i>PaCel6A</i>	7.5	57	5.9	71	
<i>PaCel6B</i>	3.3	32	4.5	40	
<i>PaCel6C</i>	3.6	23	2.7	30	
Avicel 24 h					
<i>TrCel6A</i>	246	2,119	0	2,365	7.6
<i>PaCel6A</i>	75	977	49	1,101	7.9
<i>PaCel6B</i>	43	283	8.3	335	4.7
<i>PaCel6C</i>	32	238	4.6	275	5.7
CMC 24 h					
<i>TrCel6A</i>	129	809	24	961	5.3
<i>PaCel6A</i>	12	355	69	436	4.4
<i>PaCel6B</i>	19	650	123	793	4.6
<i>PaCel6C</i>	4.4	142	16	162	7.1

^a Values were determined by HPLC.

^b For Avicel, processivity was calculated as the molar ratio (G2 - G1)/(G3 + G1), except for *PaCel6A*, where it was G2/(G1 + G3). For CMC, processivity was calculated as G2/(G3 + G1).

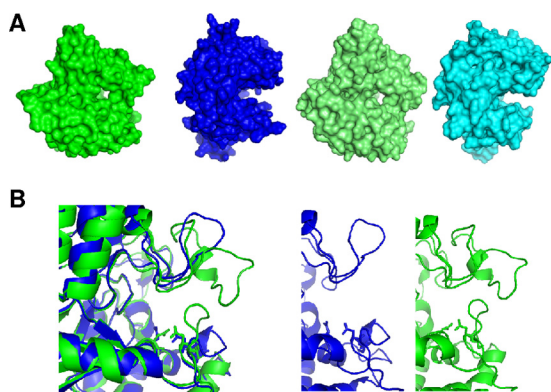


FIG 4 Structural analysis of *PaCel6*. (A) Surface representations of *PaCel6A* (left, in green), *PaCel6B* (middle left, in dark blue), *PaCel6C* (middle right, in light green), and *PaGH6D* (right, in light blue). *PaCel6A* and *PaCel6C* (green) exhibit a tunnel formation, whereas *PaCel6B* and *PaGH6D* show a cleft-shaped active site. (B) Structural alignment of the catalytic sites of *PaCel6A* (green) with *PaCel6B* (blue). The loop corresponding to amino acids 415 to 429 in *PaCel6A* leading to the formation of the tunnel shape is absent in *PaCel6B*. Models of *PaGH6* were built by using PHYRE2 and visualized with PyMOL. *PaCel6A* and *PaCel6C* models exhibited 100% confidence with *H. insolens* Cel6A (Protein Data Bank [PDB] accession numbers IOC7 and 1BVW) and *TrCel6A* (PDB accession number 1QJW), and *PaCel6B* and *PaGH6D* models showed 100% confidence with *H. insolens* Cel6B (PDB accession number 1DYS).

[37]). With time, the ongoing processive hydrolysis might then lead to proportionally less G3 and G1 formation. Glucose thus seems to originate only from false initial attack, which is confirmed by the fact that *PaCel6A*, in contrast to *TrCel6A*, cannot cleave G3.

PaCel6B and *PaCel6C* produced small amounts of G3 and G1 after 15 min. G2/G3 ratios increased after 24 h for both enzymes, whereas G2/G1 ratios remained similar. As for *TrCel6A*, this is probably due to a slow hydrolysis of G3. The ratio of $G2/(G1 + G3)$ or, for enzymes degrading G3, the ratio $(G2 - G1)/(G3 + G1)$ can be used to estimate processivity (30). Table 4 shows that processivity was lower for *PaCel6B* and *PaCel6C* than for *PaCel6A* and *TrCel6A*.

All four tested enzymes were also able to hydrolyze CMC, with the highest activity observed for *TrCel6A*, followed by *PaCel6B*. *TrCel6A* released more glucose than G3, which is in contrast to the three *P. anserina* enzymes. A possible reason for this could be that *TrCel6A* slowly cleaves G3 into glucose and G2. Alternatively, more glucose than G3 could be produced by false initial attacks. The second hypothesis would be consistent with the fact that the bulkier side chains of CMC might have more difficulty in entering the tunnel-shaped active site. *PaCel6A*, *PaCel6B*, and *PaCel6C* produced more G3 than glucose, which is in contrast to the results obtained with Avicel at 24 h. Carboxymethyl cellotriose might not be a substrate for these enzymes. On the other hand, the rather high level of production of G3 by *PaCel6A* and *PaCel6B* may indicate a higher frequency of false initial attacks due to reduced processivity of these enzymes on CMC.

Endo- or exoglucanase activity? To gain information on the active-site topology of *P. anserina* Cel6 enzymes, homology models of all four *PaCel6* enzymes were built based on known template 3D structures (Fig. 4). *PaCel6A* and *PaCel6C* display active-site tunnels, suggesting that they are processive enzymes, in good

agreement with the biochemical data that demonstrate the highest activities on crystalline cellulose. The tunnel structure could also explain the lower activity on CMC, with the substituted glucose units having more difficulty entering the narrow active site. The processivity of *PaCel6A* on Avicel is similar to that of *TrCel6A*, suggesting that the types of action of the two enzymes could be similar. However, more work is needed to elucidate the reason for the lower catalytic efficiency and specific activity of *PaCel6A*. In contrast to *PaCel6A* and *PaCel6C*, the *PaCel6B* and *PaGH6D* structures are predicted to contain a binding cleft rather than a tunnel structure. These two enzymes indeed lack 15 amino acids of the C-terminal loop (residues 406 to 420, according to *TrCel6A* numbering), generating an open topography of the active site. A similar short loop was also found for *H. insolens* Cel6B, an endoglucanase, which represents the closest characterized enzyme to *PaCel6B* and *PaGH6D*, with 54 and 82% identity, respectively. Asn182 of the N-terminal loop, which is in direct contact with Arg410 of the C-terminal loop in exoglucanases (38), is also absent in the latter two enzymes (Fig. 2). The homology model thus supports the hypothesis that these enzymes have endo-type activity. In the case of *PaCel6B*, this was already suggested by the biochemical data obtained: higher activity on CMC than on Avicel and lower processivity on Avicel were found. In addition, this enzyme was shown to cleave glucomannan and β -(1,3;1,4)-glucan, suggesting that mixed hexose polymers with different β -linkages can be accommodated in the larger active site.

DISCUSSION

Since the 1960s, *P. anserina* has been used as a model for studying fundamental biological processes, such as life cycle and fungal sexual reproduction. Because this coprophilous fungus develops on herbivore dung after zygomycetes and (hemi)cellulose-degrading ascomycetes, it is believed to have to cope with the most recalcitrant plant residues. With the current intense search for efficient biomass conversion enzymes, *P. anserina* is beginning to be investigated for its biomass-degrading capacities. The publication of its genome indeed revealed a large portfolio of lignocellulose-degrading enzymes (17), which are only starting to be characterized (27). The present study is, to our knowledge, the first one describing cellulolytic enzymes of this organism.

P. pastoris has already been used successfully to express several *P. anserina* polysaccharide-degrading enzymes (27, 39) and numerous other CAZymes (40–42). In some cases, the heterologous proteins expressed in *P. pastoris* displayed a higher degree of glycosylation than their native counterparts (43–46), and this was also observed for *TrCel6A* and *PaCel6A* in the present study. Deglycosylation evidenced the presence of both *N*- and *O*-glycans in *TrCel6A*. However, *PaCel6A* could be only marginally deglycosylated. It is possible that the large number of *O*-glycosylation sites in the linker domain leads to glycan chains that are difficult to access, which might have prevented *O*-deglycosylation. The question arises as to whether enzymatic properties are affected by this high degree of glycosylation. Whereas glycosylation of CBM and catalytic domains can alter enzyme properties, such as activity and stability, glycosylation of the linker is generally considered to protect the enzyme from protease degradation (47). Concerning *TrCel6A*, glycosylation does not seem to alter its functioning; the specific activity of the recombinant *TrCel6A* enzyme on Avicel was similar to what was already reported in the literature, even if the experimental conditions were not strictly identical: 0.016

U/mg in this study, compared to 0.027 U/mg in studies by Tomme et al. (48) and Billard et al. (49), which were obtained with the native enzyme purified from *T. reesei*. The pH and temperature profiles of recombinant *TrCel6A* were also close to those previously obtained (50). It cannot be ruled out, however, that glycosylation or the presence of tags affects the biochemical properties of one of the heterologously produced *P. anserina* enzymes which could not be compared to their wild-type counterparts.

The pH and temperature profiles for the three *PaCel6* enzymes were different from those for the recently described *P. anserina* hemicellulase enzymes (27). The GH6 enzymes studied here were generally less thermostable than the hemicellulases. In contrast, *PaCel6* enzymes were active and stable in a much larger pH range, especially at alkaline pH, which was not observed for hemicellulases. Other fungal GH6 cellulases with alkaline pH optima have been described: *H. insolens* CBH2 (Cel6A) and *Magnaporthe oryzae* Cel6A show the highest activity at pH 9, but *H. insolens* Cel6B, an endoglucanase, has a much narrower pH optimum, centered at pH 7 (26, 51).

In contrast to *PaCel6B* and *PaCel6C*, *PaCel6A* harbors a CBM1 module at its N terminus. When adsorption was measured on Avicel, about 40% of the added enzymes were bound to cellulose at 4°C after 4 h, while *PaCel6B* and *PaCel6C* did not adsorb to the substrate (not shown), suggesting that *PaCel6A*-CBM1 is a functional CBM. The missing CBM in the latter two enzymes might also be the reason for the lower specific activities observed on Avicel. However, the CBM does not seem to be important for hydrolysis of CMC (a soluble substrate), as the specific activities are not correlated with its presence.

Modeling of the three-dimensional structures corroborated experimental data suggesting an endo or exo type of hydrolytic attack of *PaCel6* enzymes. It has been known for quite some time, however, that the distinction between endo- and exoglucanases is not absolute and is sometimes hard to establish experimentally (16, 52–54). *H. insolens* Cel6A, a cellobiohydrolase, acts on crystalline cellulose ribbons in an endo-like fashion and hydrolyzes amorphous cellulose and CMC (26). However, its structure was shown to be very similar to that of *TrCel6A*, with two surface loops forming a substrate binding tunnel, characteristic of exoglucanases (38). Due to their ability to hydrolyze substituted glucose chains, such as CMC, both enzymes are therefore considered to be endo-processive cellobiohydrolases (24, 55). *PaCel6C* has a lower processivity on crystalline cellulose than does *TrCel6A* and hydrolyzes CMC nearly as well as crystalline cellulose. This suggests that this enzyme is better characterized as a processive glucanase with an even more pronounced endo character than the Cel6A enzyme of *T. reesei* or *H. insolens*.

Other examples of processive endoglucanases are known: *Agaricus bisporus* CEL3 possesses loops enclosing the active-site tunnel, but its activity profile was shown to be intermediate between typical cellobiohydrolases and endoglucanases (56). In prokaryotes, processive endoglucanases have been identified in *Saccharophagus degradans* (57), and a GH9 cellulase with both endo- and exoglucanase activities was described in *Cellulomonas fimi* (58).

The degree of endo-type activity can be related to different structural features, such as the structure of the catalytic core, which determines its binding strength (53). Binding of the substrate to the active site is known to involve several tryptophan residues (Trp135, Trp272, Trp367, and Trp269) (34). The most

important one, Trp135, binds to the glucose unit at the –2 subsite, and its binding strength was demonstrated to drive the movement of the cellulose chain leading to processive cleavage (11). All of these residues are conserved in the *P. anserina* Cel6 enzymes, but different electrostatic environments at this subsite could lead to different degrees of processivity.

Active-site loops are another factor that may affect processivity. These are thought to display some flexibility and undergo conformational changes in response to ligand binding, which could lead to occasional endo cleavage even if the enzyme presents a tunnel-like structure (38, 59, 60). The cleavage of fluorescein-labeled cellodextrins by CBHs supports the hypothesis that such conformational changes occur (61). Therefore, the degree of flexibility and length of the active-site loops may also explain the different extents of endo action observed for the *PaCel6* enzymes.

In a recent study, *P. anserina* secretomes that had been generated by induction with different substrates were tested for their capacity to supplement a *T. reesei* enzyme cocktail. By replacing half of the enzyme content with an equivalent amount of *P. anserina* enzymes, the hydrolysis yield of steam-exploded wheat straw could be increased up to 17% upon hydrolysis at 37°C. Proteomic analysis of the two secretomes that resulted in the highest gain of hydrolysis yield, namely, those obtained after induction by sugar beet pulp and Avicel, revealed the presence of the four *PaCel6* enzymes: *PaCel6A* and *PaCel6B* were induced by both substrates, whereas *PaCel6C* and *PaGH6D* were present in only one of the two secretomes (L. Poidevin, unpublished data). It cannot be confirmed at this point that the *PaCel6* enzymes are responsible for the improvement in hydrolysis yield observed. However, this finding indicates that even though a *P. anserina* secretome hydrolyzes micronized or steam-pretreated wheat straw less efficiently than a *T. reesei* enzyme cocktail (Poidevin, unpublished), *P. anserina* produces enzymes which can effectively increase hydrolysis yields when added to a classical *Trichoderma* cellulase cocktail. We are only at the beginning of the characterization of the enzymatic complex of *P. anserina*, which includes about 180 genes encoding plant cell wall-degrading enzymes (17). The presence of 15 GH5 genes, 6 GH7 genes, encoding putative cellobiohydrolases, and 33 GH61 genes is particularly intriguing. Transcriptomic and proteomic studies are now under way to determine which enzymes are produced in the presence of lignocelluloses and to gain a deeper mechanistic understanding of the cell wall-degrading arsenal of this fungus.

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